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TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371		RK OFFICE ATTORNEY'S DOCKET NUMBER 9424.170US01		
		U.S. APPLICATION NO. (If known, see 37 C.F.R. 1.5) Unknown		
NAME OF THE PARTY	INTERNATIONAL FILING DATE	PRIORITY DATE CLAIMED		
		·		
	August 29, 2000	August 30, 1999		
TITLE OF INVENTION				
COSMETICS COMPOSITION				
APPLICANT(S) FOR DO/EO/US				
VOSS, et al.				
Applicant herewith submits to the United States De	signated/Elected Office (DO/EO/US) the	following items and other information:		
[X] This express request to begin national examination until the expiration of the ap [X] A proper Demand for International Prelin	NT submission of items concerning a filin amination procedures (35 U.S.C. 371(f)) a plicable time limit set in 35 U.S.C. 371(b) hinary Examination was made by the 19th	t any time rather than delay) and PCT Articles 22 and 39(1).		
b. [X] has been transmitted by the Inte	I only if not transmitted by the Internation	iving Office (RO/US)		
b. [] have been transmitted by the	equired only if not transmitted by the Inter ne International Bureau. eer, the time limit for making such amend	rnational Bureau).		
3. [] A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).				
9. [X] An unsigned oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)).				
10. [] A translation of the annexes to the Ir (35 U.S.C. 371(c)(5)).	ternational Preliminary Examination Rep	oort under PCT Article 36		
Items 11. to 16. below concern document(s) or in 11. [] An Information Disclosure Statemer				
12. [] An assignment document for record	ing. A separate cover sheet in compliance	with 37 CFR 3.28 and 3.31 is included.		
13. [X] A FIRST preliminary amendment. [] A SECOND of SUBSEQUENT pre	liminary amendment.			
14. [] A substitute specification.				
15. [] A change of power of attorney and/o	or address letter.			
16. [X] Other items or information: Application	Data Sheet; Form PCT/IPEA/409 with A	mended Claim sheets; PCT/ISA/210		

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JC10 Rec'd PCT/PTO 1 2 FEB 2002

S/N unknown

PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant:

VOSS et al.

Docket No.:

9424.170USWO

Serial No.:

unknown

Filed:

concurrent herewith

Int'l Appln No.:

PCT/NL00/00599

Int'l Filing Date:

August 29, 2000

Title:

COSMETICS COMPOSITION

CERTIFICATE UNDER 37 CFR 1.10

'Express Mail' mailing label number: EL 669944187 US

Date of Deposit: February 12, 2002

I hereby certify that this correspondence is being deposited with the United States Postal Service 'Express Mail Post Office To Addressee' service under 37 CFR 1.10 on the date indicated above and is addressed to the Assistant Commissioner for Patents, Washington, D.C. 20231.

Name: Chris Stordahl

PRELIMINARY AMENDMENT

Box PCT Assistant Commissioner for Patents Washington, D. C. 20231

Dear Sir:

In connection with the above-identified application filed herewith, please enter the following preliminary amendment, , based on claims amended in prosecution of the international application and published in the International Preliminary Examination Report, a copy of which is enclosed herewith:

IN THE ABSTRACT

Insert the attached Abstract page into the application as the last page thereof.

IN THE SPECIFICATION

A courtesy copy of the present specification is enclosed herewith. However, the World Intellectual Property Office (WIPO) copy should be relied upon if it is already in the U.S. Patent Office.

IN THE CLAIMS

Please amend claims 3-6 and 8-16 to read as follows:

- 3. (AMENDED) Composition according to claim 1, wherein the container is a plastic container.
- 4. (AMENDED) Composition according to claim 1, wherein said composition comprises at least two surfactants and a hydrophobic compound having a HLB-value of less than 10 wherein said composition further has a viscosity according the Brookfield (23°C, Spindle TE, 5 RPM) of in the range of 5,000 to 50,000 mPas.
- 5. (AMENDED) Composition according to claim 1, comprising from 0.01 to 30 wt.% of thickener, from 1 to 15 wt.% of propellant, from 0.5 to 50 wt.% of surfactant and the balance being water and other customary body care ingredients.
- 6. (AMENDED) Composition according to claim 1, wherein the thickener is chosen from the group of gums, poly(meth)acrylates, polymers based upon aerosil-types, polysaccharides, high molecular polyethyleneglycolmono- and diesters of fatty acids, polyacrylamides, polyvinylalcohols, polyvinylpyrrolidons, esters of fatty acids with polyols, fatty alcoholethoxylates, alkyloligoglucosides and sugar-esters.
- 8. (AMENDED) Composition according to claim 1 further comprising one or more ingredients chosen from the group of pH regulating agents, oil bodies, emulsifying agents, preservatives, perfumes, moisturizing agents, UV-filters, emollients, superfatting agents, brighteners, strength improving agents, silicon agents, fats, waxes, lecithins, phospholipids, stabilizing agents, anti-bacterial agents and other bioactive agents, odor-absorbing agents, antiperspirants, antidandruff agents, film-forming agents, swelling agents, antioxidants, insect-repellents, hydrotropes, tanning agents, tyrosin inhibitors, solubilizers and colorants.

- 9. (AMENDED) Composition according to claim 1, wherein said composition comprises a fatty alcohol preferably of the formula R₁OH, R₁ being a aliphatic hydrocarbon group containing 6 to 22 carbon atoms and 0, 1, 2 or 3 double bonds.
- 10. (AMENDED) Composition according to claim 1, wherein said composition comprises a fatty alcoholalkoxylate preferably of the formula R₂O(AlkO)_mH, R₂ being an aliphatic hydrocarbon group containing 6 to 22 carbon atoms, m being an integer from 1 to 30 and AlkO being an alkyleneoxide.
- 11. (AMENDED) Composition according to claim 1, wherein said composition comprises a fatty alcoholalkoxylate of the formula R₂O(AlkO)_mH, R2 being an aliphatic hydrocarbon group containing 8 to 22 carbon atoms, m being an integer from 5 to 20 and AlkO being chosen from ethyleneoxide and propylene oxide.
- 12. (AMENDED) Composition according to claim 4, wherein the viscosity according to Brookfield (23°C, Spindle TE, 5 RPM) of said composition is in the range of 10,000 to 50,000, preferably in the range of 20,000 to 30,000 mPas.
- 13. (AMENDED) Composition according to claim 1 wherein said composition is a gel.
- 14. (AMENDED) Composition according to claim 1 wherein said composition comprises:
 - a) 0.01 to 30 % w/w of a thickener,
 - b) 0.1 to 20 % w/w of a hydrophobic compound having an HLB value of less than 10,
 - c) 0.5 to 40 % anionic surfactants,
 - d) 0.25 to 5 % amphoteric surfactants, and/or

- e) 0.5 to 40 % nonionic surfactants, and is further characterized by the composition having a viscosity of 5,000 to 50,000 mPas and the weight ratio of components c:d or c:e being in the range of 2:1 to 8:1.
- 15. (AMENDED) Container comprising a cosmetics composition according to claim1.
- 16. (AMENDED) Use of a composition according to claim 1 as a sun cream or lotion, body milk, shampoo, bathing or shower gel, ointment, deodorant, hair care product or moisturizing cream.

REMARKS

The above preliminary amendment is made to remove multiple dependencies from claims 3-6 and 8-16.

A new abstract page is supplied to conform to that appearing on the publication page of the WIPO application, but the new Abstract is typed on a separate page as required by U.S. practice.

Applicants respectfully request that the preliminary amendment described herein be entered into the record prior to calculation of the filing fee and prior to examination and consideration of the above-identified application.

If a telephone conference would be helpful in resolving any issues concerning this communication, please contact Applicants' primary attorney-of record, John J. Gresens (Reg. No. 33,112), at (612) 371.5265.

Respectfully submitted,

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Dated: February 12, 2002

John J. Gresens

JJG/pjk

ABSTRACT

The invention relates to a cosmetics composition comprising a thickener, a propellant, a surfactant and water, wherein the composition is contained in a container under a pressure of nor more than 3 bar.

MARKED-UP COPY OF CLAIMS

- 3. Composition according to claim 1 [or 2], wherein the container is a plastic container.
- 4. Composition according to [any of the claims 1-3] <u>claim 1</u>, wherein said composition comprises at least two surfactants and a hydrophobic compound having a HLB-value of less than 10 wherein said composition further has a viscosity according the Brookfield (23°C, Spindle TE, 5 RPM) of in the range of 5,000 to 50,000 mPas.
- 5. Composition according to [any of the preceding claims] <u>claim 1</u>, comprising from 0.01 to 30 wt.% of thickener, from 1 to 15 wt.% of propellant, from 0.5 to 50 wt.% of surfactant and the balance being water and other customary body care ingredients.
- 6. Composition according to [any of the preceding claims] <u>claim 1</u>, wherein the thickener is chosen from the group of gums, poly(meth)acrylates, polymers based upon aerosiltypes, polysaccharides, high molecular polyethyleneglycolmono- and diesters of fatty acids, polyacrylamides, polyvinylalcohols, polyvinylpyrrolidons, esters of fatty acids with polyols, fatty alcoholethoxylates, alkyloligoglucosides and sugar-esters.
- 8. Composition according to [any of the preceding claims] claim 1 further comprising one or more ingredients chosen from the group of pH regulating agents, oil bodies, emulsifying agents, preservatives, perfumes, moisturizing agents, UV-filters, emollients, superfatting agents, brighteners, strength improving agents, silicon agents, fats, waxes, lecithins, phospholipids, stabilizing agents, anti-bacterial agents and other bioactive agents, odor-absorbing agents, antiperspirants, antidandruff agents, film-forming agents, swelling agents, antioxidants, insect-repellents, hydrotropes, tanning agents, tyrosin inhibitors, solubilizers and colorants.

- 9. Composition according to [any of the preceding claims] <u>claim 1</u>, wherein said composition comprises a fatty alcohol preferably of the formula R₁OH, R₁ being a aliphatic hydrocarbon group containing 6 to 22 carbon atoms and 0, 1, 2 or 3 double bonds.
- 10. Composition according to [any of the preceding claims] claim 1, wherein said composition comprises a fatty alcoholalkoxylate preferably of the formula R₂O(AlkO)_mH, R₂ being an aliphatic hydrocarbon group containing 6 to 22 carbon atoms, m being an integer from 1 to 30 and AlkO being an alkyleneoxide.
- 11. Composition according to [any of the preceding claims] <u>claim 1</u>, wherein said composition comprises a fatty alcoholalkoxylate of the formula R₂O(AlkO)_mH, R2 being an aliphatic hydrocarbon group containing 8 to 22 carbon atoms, m being an integer from 5 to 20 and AlkO being chosen from ethyleneoxide and propylene oxide.
- 12. Composition according to [any of the] claim 4 [to 11], wherein the viscosity according to Brookfield (23°C, Spindle TE, 5 RPM) of said composition is in the range of 10,000 to 50,000, preferably in the range of 20,000 to 30,000 mPas.
- 13. Composition according to [any of the preceding claims] <u>claim 1</u> wherein said composition is a gel.
- 14. Composition according to [any of the preceding claims] <u>claim 1</u> wherein said composition comprises:
 - a) 0.01 to 30 % w/w of a thickener,
 - b) 0.1 to 20 % w/w of a hydrophobic compound having an HLB value of less than 10,
 - c) 0.5 to 40 % anionic surfactants,
 - d) 0.25 to 5 % amphoteric surfactants, and/or

- e) 0.5 to 40 % nonionic surfactants, and is further characterized by the composition having a viscosity of 5,000 to 50,000 mPas and the weight ratio of components c:d or c:e being in the range of 2:1 to 8:1.
- 15. Container comprising a cosmetics composition according to [any of the preceding claims] claim 1.
- 16. Use of a composition according to [any of the claims 1-14] <u>claim 1</u> as a sun cream or lotion, body milk, shampoo, bathing or shower gel, ointment, deodorant, hair care product or moisturizing cream.

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Title: Cosmetics composition

ointments, milks, aerosols, pastes and so forth.

The invention relates to a cosmetics composition and to its use.

Cosmetics compositions are nowadays available in many different forms. One product having a very distinct appearance from another product may nevertheless serve the same purpose as said other product. Examples of the wide variety that cosmetics may have include lotions, gels, creams,

The present invention seeks to provide a new form for a cosmetics composition. The objective form is a viscous composition, preferably a gel, which, after application to the skin, creates a post-application foaming effect. When the gel is contacted with the skin, it is desired that a noticeable transition takes place from a gel to a dense creamy foam. To achieve this goal, the invention provides a cosmetics composition comprising a thickener, a propellant, a surfactant and water, wherein the composition is contained in a container under a pressure of no more than 3 bar.

Surprisingly, it has been found that a propellant can be incorporated into a cosmetics composition wherein the resultant pressure is less than that of the propellant on its own. Additionally, the propellant is incorporated into the cosmetics composition substantially without affecting the stability of the composition, even if the composition has the form of a gel. When the composition is applied to the skin, the propellant is released from the gel and the user experiences a foam which is very rich, creamy and long-lasting. It has further been found that the composition is non-flammable and as such is associated with a reduced risk for consumers with respect to fire and explosion hazards.

It is to be noted that cosmetics compositions in the form of a gel and comprising a propellant are known per se. In the field of shaving creams, gels are marketed which, due to the presence of a propellant, convert into a foaming layer or lather when brought into contact with the skin. More recently, this technology has been modified to allow shower foam products to be marketed. These compositions are generally packaged in containers having two compartments, e.g. a bag made of a laminated material suspended inside an aerosol can. Moreover, the amount of propellant needed in these compositions is relatively high. Hence, shaving gels are packed into pressurized containers wherein the pressure usually is as high as 8 bar or

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more. Consequently, the container needs to be particularly strong and is therefore made of metal.

A composition according to the invention does not require such high amounts of propellant. As has been mentioned, the pressure in a container in which the composition is contained may be as low as 3 bar or less. Preferably, said pressure is lower than 2 bar. Due to this low pressure, the present composition may advantageously be packed in a plastic container, e.g. a polyethylene or polypropylene container, which is economically much more attractive than package in a metal container. Furthermore, a composition according to the invention has been found to have a lower pressure at elevated temperatures than expected. This feature is advantageous in that it can often not be avoided that the product needs to be stored at elevated temperatures for a certain period of time. It is to be noted that the pressure refers to an absolute pressure. Furthermore, the pressure will be in excess of atmospheric pressure, i.e. an overpressure of at least 0.1 bar.

It is preferred that the present composition has the form of a gel. Accordingly, the composition comprises a thickener, which is preferably present in an amount of from 0.01 to 30 wt.%, more preferably from 0.5 to 10 wt.%, even more preferably from 1 to 3 wt.%, based on the weight of the composition. Suitable thickeners are chosen for the compatibility with the propellant and for their capability to provide a stable viscous product, more in particular a stable gel. For many applications a suitable viscosity is a viscosity of 5,000 to 50,000, preferably 10,000 to 50,000, and more preferably 20,000 to 30,000 mPas, measured as Brookfield viscosity (23°C, spindle TE, 5 Rpm), e.g. to employ in cleansing gels, face cleansing gels, face care gels, gels for masks, and similar compositions which are commonly used in body care.

Examples of thickeners to be used in compositions according to the present invention include side chain modified polymers based on aerosil-types (hydrophilic silicic acids), polysaccharides such as xanthan gum, guar-guar, agar-agar, alginates and tyloses, carboxymethylcellulose, hydroxyethylcellulose, high molecular polyethylene glycol mono- and diesters of fatty acids, polyacrylates (e.g. Carbopole® of Goodrich or Synthalene® of Sigma), polyacrylamide, polyvinyl alcohols and polyvinylpyrrolidon. Particularly, thickeners with associate action, such as fatty acid glycerides, esters of fatty acids with polyols such as pentaerythrit or trimethylolpropane,

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fatty alcohol ethoxylates, optionally with EO-homologue distribution, alkyloligoglucosides and sugar esters may be used.

In accordance with the invention, highly preferred thickeners are gums and poly(meth)acrylates, such as polyacrylic acid. Particularly stable gels have been obtained using xanthan gum as the thickener. The presence of a molecular network is highly beneficial to the stability of the gel and can be demonstrated in rheological measurements.

A further important component of the present composition is a propellant. Advantageously, the propellant is chosen for its physico-chemical properties and the character of the foam texture produced on application to skin. A further consideration on which the choice for a suitable propellant may be based is its environmentally friendly character. Possible propellants are hydrocarbons, such as alkanes having from 1 to 12 carbon atoms. Preferred propellants are hydrocarbons with 4 to 7 carbon atoms, such as linear or branched alkanes, including cyclic ones. Particularly good results have been obtained using isopentane as a propellant. The propellant is preferably present in an amount suitable to achieve the desired pressure of the composition in a container in which it is packed. Suitable amounts range from 0.1 to 20 wt.%, preferably from 1 to 15 wt.%, more preferably from 4 to 8 wt.%, based on the weight of the composition.

Furthermore it has been found unexpectedly that alkanes in combination with thickeners, can yield highly viscous mixtures, of a gel-like nature. In cosmetic or pharmaceutical applications, these gel-like mixtures demonstrate a high stability, also when the temperature is decreased. Another advantage of compositions according to the invention may be, the pleasant sensation to the skin, compositions according to the invention may give rise to.

The composition further comprises one or more surfactants. Preferably, the surfactant or surfactants are foaming and skin friendly. Possible surfactants include anionic, nonionic and/or amphoteric surfactants. Typical examples of anionic surfactants include soaps, alkylbenzol sulfonates, alkane sulfonates, olefin sulfonates, alkyl ether sulfonates, glycerine ether sulfonates, α-methylester sulfonates, sulfofatty acids, alkyl sulfates, fatty alcohol ether sulfates, glycerine ether sulfates, fatty acid ether sulfates, hydroxy mixed ether sulfates, monoglyceride (ether) sulfates, fatty acid amide(ether) sulfates, mono- and dialkylsulfosuccinates, mono- and dialkyl sulfosuccinamates, sulfotriglycerides, amide soaps, ether carboxylic acids and

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salts thereof, fatty acid isothionates, fatty acid sarcosinates, fatty acid taurides, N-acylamino acids such as acylactylate, acyltartrate, acylglutamate and acylaspartate, alkyloligoglucoside sulfates, protein fatty acid condensates and alkyl (ether) phosphates. Typical examples of nonionic surfactants include fatty alcoholpolyglycol ethers, alkylphenolpolyglycol ethers, fatty acid polyglycol esters, fatty acid amidpolyglycol ethers, fatty aminpolyglycol ethers, alkoxylated triglycerides, mixed ethers, optionally partially oxidized alk(en)yloligoglycosides or glucoronic acid derivates, fatty acid-Nalkylclycamides, proteinhydrolysates, polyol fatty acid esters, sugar esters, sorbitan esters, polysorbates and amino-oxides. Typical examples of amphoteric or zwitterionic surfactants include alkyl betains, alkylamido betains, aminopropionates, aminoglycinates, imidazolinium betains and sulfo betains. Further reference, e.g. concerning the preparation of these compounds, may be made to J. Falbe (ed.), "Surfactants in Consumer Products", Springer Verlag. Berlin. 1987, p. 54-124 or J. Falbe (ed.), "Katalysatoren, Tenside und Mineralöladditive", Thieme Verlag, Stuttgart, 1978, p. 123-217. Examples of preferred surfactants include polysorbate 20 or 40, coco glucoside, lauryl glucoside, decyl glucoside, lauryl sulfates such as ammonium, sodium, magnesium, MEA, TEA, or Mipa lauryl sulfate, cocamidopropyl betain, and sodium alkyl sulfosuccinates. The surfactant is preferably present in an amount of from 0.5 to 50 wt.%, more preferably from 2 to 20 wt.% and most preferably from 8 to 13 wt.%, based on the weight of the composition.

It is possible to use a combination of ionic surfactants and amphoteric or non-ionic surfactants. Preferably the ionic surfactant is an anionic surfactant in such combinations. Typically the concentration ranges of such a composition comprising ionic surfactants and amphoteric or non-ionic surfactants are: 0.01 to 30 wt.% of a thickener, 0.1 to 20 wt.% of a hydrophobic compound having an HLB (hydrophilic/lipophilic balance) value of less than 10, 0.5 to 40 wt.% anionic surfactants, 0.25 to 5 wt.% amphoteric surfactants and/or 0.5 to 40 wt.% nonionic surfactants. Furthermore such combinations are preferably used in a ionic surfactant to amphoteric/non-ionic surfactant of in the range of 2:1 to 8:1 wt. to wt, more preferably in the range of 4:1 to 6:1 wt. to wt.

The present compositions may further comprise fatty alcohols, by which primary aliphatic alcohols of the formula R₁OH are meant, in which R₁

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is an aliphatic hydrocarbon group containing 6 to 22, preferably 10 to 18 carbon atoms and 0, 1, 2 or 3 double bonds. Typical examples are capron alcohol, capryl alcohol, 2-ethylhexyl alcohol, caprin alcohol, lauryl alcohol, isotridecyl alcohol, myristyl alcohol, cetyl alcohol, palmoleyl alcohol, stearyl alcohol, isostearyl alcohol, oleyl alcohol, elaidyl alcohol, petroselinyl alcohol, linolyl alcohol, linolenyl alcohol, elaeostearyl alcohol, arachyl alcohol, gadoleyl alcohol, behenyl alcohol, erucyl alcohol, and brassidyl alcohol and mixtures thereof. These compounds may be present in amounts of 0.1 to 20, preferably 0.5 to 10 wt.% based on the weight of the composition.

The present compositions may further comprise fatty alcohol ethoxylates, which may have the formula R₂O(AlkO)_mH, in which R₂ is an aliphatic hydrocarbon group containing 6 to 22, preferably 10 to 18 carbon atoms and 0, 1, 2 or 3 double bonds, m is an integer from 1 to 30, preferably 5 to 20, more preferably 10 to 15, and AlkO is an alkylene oxide. AlkO may be chosen from ethylene oxide, propylene oxide and/or butylene oxide. These compounds may be present in amounts of 0.1 to 20, preferably 0.5 to 10 wt.% based on the weight of the composition.

Depending on the envisaged purpose of the composition, one or more other ingredients may be present. Examples of such ingredients include pH regulating agents, oil bodies, emulsifying agents, preservatives, perfumes, moisturizing agents, UV-filters, emollients, superfatting agents, brighteners, strength improving agents, silicon agents, fats, waxes, lecithins, phospholipids, stabilizing agents, anti-bacterial agents and other bioactive agents, odorabsorbing agents, antiperspirants, antidandruff agents, film-forming agents, swelling agents, antioxidants, insect-repellents, hydrotropes, tanning agents, tyrosin inhibitors, solubilizers and colorants. The composition may further comprise a conventional cosmetics base, such as water, oil, ointment etc..

The composition is preferably formulated to be a gel. In a preferred embodiment the gel transforms upon dispensing from a container into a soft and foamy mousse which cleans in a soft and silky manner.

The pH of the composition is preferably regulated to be close to the pH of the skin itself. Accordingly, the pH of the composition is preferably slightly acidic to slightly alkaline, e.g. in the range of 5 to 8. An example of a suitable pH regulating agent is citric acid. The skilled person will be aware of numerous suitable pH regulating agents that may be employed in the present

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type of compositions. The amount of the pH regulating agent present is of course adjusted so that the desired pH is achieved.

Examples of suitable oil bodies are guarbeta-alcohols based upon fatty alcohols containing 6 to 18 or preferably 8 to 10 carbon atoms, esters of linear C₆-C₂₂ fatty acids with linear C₆-C₂₂ fatty alcohols, esters of branched C₆-C₁₃ Carboxylic acids with linear C₆-C₂₂ fatty alcohols, such as myristylmyristate, myristyl-palmitate, myristyl-stearate, myristyl-isostearate, myristyl-oleate, myristyl-behenate, myristyl-erucate, cetyl-myristate, cetylpalmitate, cetyl-stearate, cetyl-isostearate, cetyl-oleate, cetyl-behenate, cetylerucate, stearyl-myristate, stearyl-palmitate, stearyl-stearate, stearylisostearate, stearyl-oleate, stearyl-behenate, stearyl-erucate, isostearylmyristate, isostearyl-palmitate, isostearyl-stearate, isostearyl-isostearate, isostearyl-oleate, isostearyl-behenate, isostearyl-oleate, oleyl-myristate, oleylpalmitate, oleyl-stearate, oleyl-isostearate, oleyl-oleate, oleyl-behenate, oleylerucate, behenyl-myristate, behenyl-palmitate, behenyl-stearate, behenylisostearate, behenyl-oleate, behenyl-behenate, behenyl-erucate, erucylmyristate, erucyl-palmitate, erucyl-stearate, erucyl-isostearate, erucyl-oleate, erucyl-behenate and erucyl-erucate. Other examples are esters of linear C6-C22 fatty acids with branched alcohols, in particular with 2-ethylhexanol, esters of carboxylic acids with linear or branched C₆-C₂₂ fatty alcohols, in particular dioctylmalates, esters of linear and/or branched fatty acids with multivalent alcohols, such as propyleneglycol, dimerdiols or trimertriols, and/or guarbetaalcohols, triglycerides based upon C6-C10 fatty acids, liquid mono-/di-/triglyceride mixtures based upon C₆-C₁₈ fatty acids, esters of C₆-C₂₂ fatty alcohols and/or guarbeta-alcohols with aromatic carboxylic acids, in particular benzoic acid, esters of C2-C12 dicarboxylic acids with linear or branched alcohols having 1 to 22 carbon atoms or polyols with 2 to 10 carbon atoms and 2 to 6 hydroxyl groups, oils of vegetable origin, branched primary alcohols, substituted cyclohexanes, linear and branched C6-C22 fatty alcohol carbonates, guarbetacarbonates, esters of benzoic acid and linear and/or branched C₆-C₂₂ alcohols, such as Finsolv® TN, linear or branched, symmetric or asymmetric dialkylethers having 6 to 22 carbon atoms per alkyl group, ring opening products of epoxy fatty acid esters and polyols, silicon-oils and/or aliphatic respectively naphtalenic hydrocarbons, such as squalane, squalene or dialkylcyclohexanes.

An emollient or moisturizing agent may be present in order to improve the ease of application of the composition and the final skin feel the user experiences. Examples of suitable emollients or moisturizing agents include glycerin, propenylglycol, PEG 7 glyceryl cocoate, PEG 6 caprylic or capric glycerides, glyceryl oleate and lipids in general, such as paraffin oil or polar oils. An emollient or moisturizing agent is preferably present in an amount ranging from 0.5 to 15 wt.%, based on the weight of the composition.

Suitable emulsifying agents are for example non-ionic surfactants such as:

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- reaction products of 2 to 30 mole ethylene oxide and/or 0 to 5 mole propylene oxide with linear fatty alcohols having 8 to 22 carbon atoms, with fatty acids having 12 to 22 carbon atoms, with alkylphenols having 8 to 15 carbon atoms in the alkyl group and also with alkylamines having 8 to 22 carbon atoms in the alkyl group;
- alkyl and/or alkenyl-oligoglycosides having 8 to 22 carbon atoms in alk(en)yl group and ethoxylated analogs thereof;
- reaction products of 1 to 15 mole ethylene oxide and castor oil and/or fixated castor oil;
- reaction products of 15 to 60 mole ethylene oxide and castor oil and/or fixated ricinoleic oil;
- partial-esters of glycerin and/or sorbitane with unsaturated, linear or saturated, branched fatty acids having 12 to 22 carbon atoms and/or hydroxycarboxylic acids having 3 to 18 carbon atoms, as well as adducts thereof with 1 to 30 mole ethylene oxide;
- partial-esters of polyglycerin (average self-condensation degree 2 to 8), polyethyleneglycol (molecular weight of 400 to 5000), trimethylolpropane, pentaerythrite, sugar-alcohols, such as sorbitol, alkylglucosides, such as methylglucoside, butylglucoside, laurylglucoside, as well as polyglucosides, such as cellulose, with saturated and/or unsaturated, linear or branched fatty acids having 12 to 22 carbon atoms and/or hydroxycarboxylic acids having 3 to 18 carbon atoms, as well as adducts thereof with 1 to 30 mole ethylene oxide;

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- mixed-esters of pentaerythrite, fatty acids, citric acid and fatty alcohols in accordance with DE 1165574 PS and/or mixed-esters of fatty acids having 6 to 22 carbon atoms, methylglucose and polyols, preferably glycerin or polyglycerin;
- mono-, di- and trialkylphosphates, as well as mono-, di- and/or tri-PEG-alkylphosphates and salts thereof;
- wool-wax alcohols;
- polysiloxan-polyalkyl-polyether-copolymers respectively derivatives thereof;
- polyalkelyneglycols;
- glycerincarbonates;

The reaction products of ethylene oxide and/or propylene oxide with fatty alcohols, fatty acids, alkylphenols or ricinolic oils are commercially obtainable products. They are available as homologous mixtures, of which the average alkoxylation degree is in accordance with the mass ratio of ethylene oxide and/or propylene oxide and substrate, with which the reaction takes place. C₁₂-C₁₈ fatty acid mono- and diesters of reaction products of ethylene oxide and glycerin are known in relation to cosmetic compositions from DE-2024051.

Typical examples of suitable partial-glycerides are monoglyceride-hydroxy-stearinate, diglyceridehydroxy-stearinate, monoglyceride-isostearinate, monoglyceride-oleiate, diglyceride-oleiate, diglyceride-linoleate, monoglyceride-linoleate, monoglyceride-linoleate, diglyceride-linoleate, diglyceride-linoleate, diglyceride-linoleate, monoglyceride-erucate, diglyceride-erucate, monoglyceride-tartrate, diglyceride-tartrate, monoglyceride-citrate, diglyceride-citrate, monoglyceride-malate, diglyceridemalate, as well as technical mixtures thereof, which may still contain small amounts of triglyceride, depending upon the production process. Reaction products of 1 to 30, preferably 5 to 10 mole ethylene oxide with aforementioned partial-glycerides are suitable too.

Examples of suitable sorbitol-esters are sorbitol-monoisostearate, sorbitol-sesquiisostearate, sorbitol-diisostearate, sorbitol-triisostearate, sorbitol-monooleate, sorbitol-sesquioleate, sorbitol-dioleate, sorbitol-trioleate, sorbitol-monoerucate, sorbitol-sesquierucate, sorbitol-dierucate, sorbitol-trierucate, sorbitol-monoricinoleate, sorbitol-sesquiricinoleate, sorbitol-di-

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ricinoleate, sorbitol-triricinoleate, sorbitol-monohydroxystearate, sorbitol-sesquihydroxy-stearate, sorbitol-dihydroxystearate, sorbitol-trihydroxy-stearate, sorbitol-monotartrate, sorbitol-sesquitartrate, sorbitol-ditartrate, sorbitol-tritartrate, sorbitol-monocitrate, sorbitol-sesquicitrate, sorbitol-dicitrate, sorbitol-tricitrate, sorbitol-monomaleate, sorbitol-sesquimaleate, sorbitol-dimaleate, sorbitol-trimaleate as well as technical mixtures thereof. Reaction products of 1 to 30, preferably 5 to 10 mole ethylene oxide with said sorbitol esters are suitable too.

Typical examples of suitable polyglycerin esters are polyglyceryl-2 dipolyhydroxy-stearate (Dehymuls® PGPH), polyglycerin-3-diisostearate (Lameform® TGI), polyglyceryl-4 isostearate (Isolan® Gl 34), polyglyceryl-3 oleate, diisostearoyl polyglyceryl-3 diisostearate (Isolan® PDI), polyglyceryl-3 methylglucose distearate (Tego Care® 450), polyglyceryl-3 beeswax (Cera Bellina®), polyglyceryl-4 caprate (Polyglycerol Caprate T20 10/90), polyglyceryl-3 cetyl ether (Chimexane® NL), polyglyceryl-3 distearate (Cremophor® GS 32) and polyglyceryl polyricin-oleate (Admul® WOL 1403) polyglyceryl dimerateisostearate, as well as, mixtures thereof.

Further examples of suitable polyol esters are possibly with 1 to 30 mole ethylene oxide derivatized mono-, di- and triesters of trimethylolpropane or pentaerythrite with lauric acid, cocinic acid, palmic acid, talcum-oil acid, palmic acid, stearic acid, oleic acid, behenic acid and the like.

Cationic surfactants may also be suitable emulsifying agents.

Preferred cationic surfactant are quaternary esters, in particular quaternary methyl-di-fatty acid- triethanol-amine-ester salts.

Superfatting agents may be compounds such as lanolin or lecithin as well as polyethoxyated or acylated lanolin- and lecithin derivatives, polyol fatty acid esters, monoglycerides and fatty acid alkanolamides, wherein fatty acid alkanolamides also tend to stabilizing the foam.

Brighteners may for example be selected from: alkylene glycolesters, in particular ethylene glycoldistearate, fatty acid alkanolamide, in particular cocinic acid diethanolamide; partial-glycerides, especially monoglyceridestearate; esters of multivalent, possibly hydroxyl substituted carboxylic acids with fatty alcohols having 6 to 22 carbon atoms, in particular esters with long chains of tartaric acid; fatty compounds, such as fatty acids, fatty ketones, fatty aldehydes, fatty ethers and fatty carbonates, which have a total of at least 24 carbon atoms, in particular laurin and distearyl ether; fatty

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acids such as stearic acids, hydroxystearic acids or behenic acids, ring opening products of olefinepoxides having 12 to 22 carbon atoms with fatty alcohols having 12 to 22 carbon atoms and/or polyols having 2 to 15 carbon atoms and 2 to 10 hydroxyl groups, as well as mixtures thereof.

As strength improving agents, particularly suitable groups of compounds are e.g. C₁₂-C₂₂ fatty alcohols or hydroxy- fatty alcohols, preferably having 16 to 18 carbon atoms. Partial-glycerides, fatty acids or hydroxy fatty acids are also examples of suitable strength improving agents. Preferred is a combination of these compounds with alkyloligoglucosides and/or fatty acids-N-methylglucamides of equal chain length and/or polyglycerinpoly-12-hydroxystearates.

Examples of suitable silicon compounds are dimethylpolysiloxanes, methylphenylpolysiloxanes, cyclic silicons, as well as amino-, fatty acid-, alcohol-, epoxy-, fluoro-, glycoside-, and/or alkylated silicon compounds, which may either be in a liquid phase or a resin phase, at room temperature. Other examples are dimethicones, in particular mixtures of dimethicones having an average chain length of 200 to 300 dimethylsiloxane moieties and hydrated silicates. A detailed overview of suitable volatile silicons can be found in Todd et al., Cosm. Toil. 91, 27 (1976).

Typical examples of fats are tri-glycerides. Suitable waxes are for example natural waxes, such as candililla wax, carnauba wax, Japan wax, esparto grass wax, ceric wax, guaruma wax, rice bran wax, sugar cane wax, orycury wax, montan wax, bees wax, shellac wax, walrat, lanolin (wool wax), tail root fat, ceresin, ozocerite (earth wax), petrolatum, paraffin wax, micro waxes, chemically modified waxes (hard waxes) such as montan ester waxes, sasol waxes, hydrated jujube waxes, as well as synthetic waxes, such as polyalkylene waxes and polyethyleneglycol waxes. In addition to fats, certain compounds that are similar to fats may be added, such as lecithins and phospholipids. With lecithins, the person skilled in the art means those glycero-phospholipids, which can be formed by esterification of fatty acids, glycerin, phosphoric acids and choline. In the art, lecithins are therefor also often referred to as phospatidylcholine (PC) and can be characterized by the following general formula:

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$$\begin{array}{c} CH_2OCOR \\ RCOO\text{-}CH \\ O \\ CH_3 \\ CH_2O\text{-}P\text{-}OCH_2CH_2\text{-}N^+\text{-}CH_3 \\ O^- \\ CH_3 \end{array}$$

wherein R typically represents linear aliphatic hydrocarbon moieties having 15 to 17 carbon atoms and up to 4 cis-double bonds. Examples of natural lecithins are compounds from the group of Cephalins, which are also referred to as phosphatide acids, and derivatives of 1,2-diacyl-sn-glycerin-3-phosphoric acids.

Examples of suitable phospholipids are mono-esters and, preferably, di-esters of phosphoric acids and glycerin (i.e. glycerin phosphates), which are generally regarded as fatty substances. Sphingosines or better sphingolipids are other examples of suitable additives.

Metal salts of fatty acids, such as magnesium- aluminum- and/or zinc stearate respectively -ricinoleate can be employed as stabilizing agents.

Suitable biologically active additives include tocopherol, tocopherol acetate, tocopherol palmitate, ascorbic acid, desoxyribonucleinic acid, retinol, bisabolol, allantoin, phytantriol, panthenol, AHA acids, amino acids, ceramides, pseudoceramides, essential oils, plant extracts and vitamin C complexes.

Cosmetic anti bacterial agents are active against the development of body odors. Body odors develop due to the activity of dermal bacteria on apocryne perspiration, during which unpleasantly smelling metabolites are formed.

Suitable anti bacterial agents include germination inhibiting compounds which are in principle active against all gram positive bacteria, such as 4-hydroxybenzoic acid and salts plus esters thereof, N-(4-chlorophenyl)-N'-(3,4 dichlorophenyl)urea, 2,4,4'-trichloro-2'-hydroxy-diphenyl ether (Triclosan), 4-chloro-3,5-dimethylphenol, 2,2'-methylene-bis(6-brom-4-chlorophenol), 3-methyl-4-(1-methylethyl)phenol, 2-benzyl-4-chlorophenol, 3-(4-chlorophenoxy)-1,2-propandiol, 3-iodo-propinylbutyl carbamate chlorohexidin, 3,4,4'-trichloro carbanilide (TTC), antibacterial fragrant compounds, thymol, thymian oil, eugenol, clove oil, menthol, mint oil, fernesol,

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phenoxyethanol, glycerin-monolaurate (GML), diglycerin-monocaprinate (DMC), salicylic acid-N-alkylamide such as salicylic acid-n-octylamide or salicylic acid-n-decylamide. Furthermore enzyme inhibitors can be used to help to prevent the production of undesired body odors. Esterase inhibitors, for example, are suitable for this purpose in compositions according to the invention. Preferred enzyme inhibiting agents are trialkylcitrates, such as trimethylcitrate, tripropylcitrate, triisopropylcitrate, tributylcitrate and in particular triethylcitrate (Hydragen® Cat, Henkel KGaA, Dusseldorf/FRG). The compounds inhibit the enzyme activity and reduce the formation of odorous compounds. Other suitable esterase inhibiting compounds are for example sterolsulphates or -phosphates, such as lanosterin-, cholesterin-, campesterin-, stigmasterin and sitosterinsulphate respectively -phosphate. Dicarboxylic acids and esters thereof, such as glutaric acids, glutaric acid mono-ethylesters, adipinic acid, adipinic acid monoethylester, adipinic acid diethylester, malonic acids and malonic acid diethyl ester, hydroxycarboxylic acids and esters thereof such as citric acid, malic acid, tartaric acid or tartaric acid diethylester as well as zinc glycinate.

In addition odor-absorbing agents may be used to suppress the formation of a undesired scent. Suitable compounds decrease the partial pressure of the single components and as such decrease the velocity of spreading. It is important that perfume compositions are not absorbed significantly. Odor-absorbing agents are normally not directly active against bacteria. They comprise for example as primary component a complex of a zinc salt of ricinoleic acid or special fragrance-neutral perfume compounds, known to the skilled professional as fixatives. Examples of these fixatives are labdanum extracts respectively Styrax or certain abietic acid derivatives. Furthermore perfume compounds, including fragrant oils, may serve as masking agents and also may give a typical fragrant character to compositions. Examples of fragrant oils are mixtures of natural and synthetic fragrant compounds. Examples of natural fragrant compounds are extract of flowers, stems, leafs, fruits, fruit skin, fruit peel, rots, woods, herbs, grasses, needles and branches, as well as resins and balms. Furthermore materials of animal origin are suitable, such as civet or castoreum. Typical synthetic fragrant compounds are esters, ethers, aldehydes, ketones, alcohols and hydrocarbons. Examples of fragrant esters as benzylacetate, p-tertbutylcyclohexylacetate linalyl acetate, phenylethylacetate, benzylbenzoate,

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benzylformiate, allylcyclohexyl-proprionate, styrallylproprionate and benzylsalicytate. An example of a suitable ether is benzylethyl ether. Examples of suitable aldehydes are linear alkanals having 8-18 carbon atoms, citral, citronellal, citronellyloxyacetyl aldehyde, cyclamen aldehyde, hydroxycitronellal, lillial and bourgeonal. Examples suitable ketones are ionones and methylcedrylketone. Examples of suitable alcohols are anethol, citronellol, eugenol, isoeugenol, geraniol, linalool, phenylethyl alcohol and terpineol and examples, of the hydrocarbons mainly terpenes and balms. Preferably mixtures of different fragrant compounds, resulting in a pleasant aroma, are employed. Also commonly used aromatic compounds of the group of 10 etheric oils of low volatility, are suitable perfume oils. Examples of these are sage oil, camille oil, clove oil, balm mint oil, mint oil, cinnamon oil, lindenblossom oil, juniper oil, vetiver oil, olibanum oil, galbanum oil, labdanum oil and layender oil. Preferred are bergamot oil, dihydromyrcenol, lilial, lyral, citronellol, phenylethyl alcohol, a-hexylcinnamon aldhyde, geraniol, 15 benzylacetone, cyclamen aldehyde, linalool, boisambrene forte, ambroxan, indol, hedione, sandelice, citron-oil, mandarin-oil, orange-oil, allylamylglycolate, cyclovertal, lavender-oil, muscate sage-oil, β-damascone, geranium-oil bourbon, cyclohexylsalicylate, vertofix coeur, iso-E-super, fixolide 20 NP, evemyl, iraldein gamma, phenyl-acetic acid, geranyl acetate, benzyl acetate, rose-oxide, romilate, irotyl and floramate, either employed alone or in a mixture.

Anti-perspirants reduce the formation of perspiration by influencing the activity of exocryne perspiratory glands, and as such help to prevent wetting of arm pits as well as the formation of body odors. Aqueous or non aqueous compositions of Anti-perspirants typically comprise the following ingredients:

- astringent agents;
- oil compounds;
- non ionic emulsifying agents;
- co-emulsifying agents;
- strength improving agents;
- aiding compounds such as thickening agents or complexing agents; and/or

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• non-aqueous solvents such as ethanol, propyleneglycol; and/or glycerin.

Examples of astringent agents are in particular all salts of aluminum, zirconium and zinc, such as aluminumchloride, aluminumchlorohydrate, aluminumdichlorohydrate, aluminumsesquichlorohydrate, and complexes thereof, e.g. with propyleneglycol-1,2. Aluminumhydroxyallantoinate, aluminumchloridetartrate, aluminumzirconium-trichlorohydrate, aluminum-zirconium-tetrachlorohydrate, aluminum-zirconium-pentachlorohydrate and complexes thereof, for example with amino acids such as glycin. In addition anti-perspirants may comprise the usual oil soluble and water soluble aiding agents in lower concentrations. Examples of oil soluble aiding agents are:

- infection inhibiting, skin protecting or fragrant etheric oils;
- synthetic skin protecting agents; and/or
- oil soluble perfume oils.

Suitable antidandruff agents are for example Octopirox® (1hydroxy-4-methyl-6-(2,4,4-trimethylpentyl)-2-(1H)-pyridon-monoethanolamine
salt, Babypival, Pirocton Olamin, Ketoconazol®, (4-acetyl-1-{-4-[2-(2,4-dichlorophenyl)r-2-(1H-imidazol-1-ylmethyl)-1,3-dioxylane-c-4-ylmethoxyphenyl}
piperazin, selenedisulfide, sulfur colloidals, sulfurpolyethyleneglycolsorbitolmonooleate, sulfurricin-polyethoxylate, sulfurtar destillate, salicylic acid (in
particular in combination with hexachlorophen), undexylene acid monoethanolamide sulfosuccinate Na-salt, Lamepon® UD (protein-undecylene acid
condensate, zincpyrethione, aluminumpyrethione and
magnesiumpyrethione/dipyrethione-magnesiumsulfate.

Usual film-forming agents include chitosan, micro crystalline chitosan, quaternary chitosan, polyvinylpyrolidon, vinylpyrolidon-vinylacetate-copolymerisate, polymers of acrylic acid, quaternary cellulose derivatives, collagen, hyaluronic acid, respectively salts thereof and similar compounds.

Suitable swelling agents for aqueous phases include montmorillonites, clay mineral compounds, pemules, as well as alkylated carbopoltypes (Goodrich). Furthermore, polymers suitable as swelling agents can be found in the overview by R. Lochhead in cosm. Toil. 108, 95 (1993).

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Suitable UV-filters are for example compounds - liquid or crystalline at room temperature - that are capable of absorbing ultraviolet radiation and of releasing the absorbed energy in the form of electromagnetic radiation of a longer wavelength, e.g. in the form of infra red radiation. UVB filters may be oil soluble or water soluble. Examples of oil soluble compounds are:

- 3-benzylidencamphor respectively 3-benzylidennorcamphor and derivatives thereof, such as 3-(4-mathylbenzyliden)camphor as further described in EP 0693471;
- 4-aminobenzonic acid derivatives, preferably 4-dimethylaminobenzoic acid-2-ethylhexyl ester, 4-(dimethylamino)benzoic acid-2octyl ester and 4-dimethylamino)benzoic acid amyl ester;
- esters of cinnamon acid, preferably 4-methoxy-cinnamon-2ethylhexyl ester, 4-methoxy-cinnamon acid propyl ester, 4methoxy-cinnamon acid isoamyl ester 2-cyano-3,3-phenyl cinnamon acid-2-ethylhexyl ester (octocrylene);
- esters of salicylic acid, preferably salicylic acid-2-ethylhexyl ester, salicylic-4-isopropylbenzyl ester, salicylic acid homomenthyl ester;
- derivatives of benzophenones, preferably 2-hydroxy-4methoxybenzophenone, 2-hydroxy-4-methoxy-4'methylbenzophenone, 2,2'di-hydroxy-4-methoxybenzophenone;
- esters of benzalmalon acid, preferably 4-methoxybenzamalon acid di-2-ethylhexyl ester;
- triazin derivatives, e.g. 2,4,6-trianilino-(p-carbo-2'-ethyl-1'-hexyloxy)-1,3,5-triazin and octyl triazon, as described in EP 0 818 450 A1 or dioctyl butamido triazone (Uvasorb® HEB);
- propane-1,3-dione, e.g. 1-(4-tert butylphenyl)-3-(4'methoxyphenyl)propane-1,3-dione;
- ketotricyclo(5.2.1.0)decane-derivatives as described in EP 0 694 521 B1.

Examples of water soluble UV-filters are:

 2-phenylbenzimidazol-5-sulfonic acid and alkali-, earth alkali-, ammonium-, alkylammonium-, alkanolammonium- and glucammonium salts thereof;

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- sulfonic acid derivatives of benzophenon, preferably 2-hydroxy-4-methoxybenzophenone-5-sulfonic acid and salts thereof;
- sulfonic acid derivatives of 3-benzylidencamphors, e.g. 4-(2-oxo-3-bornylidenemethyl)benzol-sulfonic acid and 2-methyl-5-(2 oxo-3-bornylidene)sulfonic acid and salts thereof.

Typical examples of UV-A filters are derivatives of benzoylmethane, such as 1-(4'-tert.butylphenyl)-3-(4'-methoxyphenyl)-propane-1,3-dione as well as enamine compounds as described in DE 19712033 A1 (BASF). Naturally it is possible to employ mixtures of UV-A and UV-B filters. In addition to the already mentioned soluble compounds, non-soluble sun-screen pigments may also be employed. In particular small dispersed metal oxide particles and metal salts, such as zinc oxide, titanium dioxide, oxides of respectively iron, zirconium, silicium, manganese, aluminum and cerium, as well as mixtures thereof, the salts of silicates (talcum), barium sulfate and zinc stearate. The oxides and salts are employed in compositions for skin care or skin protective emulsions and in decorative cosmetics. The particles should have an average diameter of less than 100 nm, preferably between 5 and 50 nm, more preferably between 15 and 30 nm. The particles may have a spherical, ellipsoidal or other shape. Optionally the surfaces of pigments may have been treated, i.e. by hydrophilization or hydrophobization. Typical examples are coated titanium dioxide, such as titanium dioxide T 805 (Degussa) or Eusolex® T2000 (Merck). Typical examples of hydrophobic coating agents are silicones. and particularly trialkoxyoctylsilanes or Simethicones. So called micro- or nanopigments are preferably employed in sun screen compositions. Preferably micronized zinc oxide is used. Further examples of suitable UV-filters can be found in the overview of P. Finkel in SÖFW-Journal 122, 543 (1996).

In addition to the groups of primary light protective agents, as mentioned above, it is also possible to use secondary light protective agents of the group of antioxidants which can stop photochemical reaction chains. These photochemical reactions are induced by UV-radiation as it enters the skin. Typical examples of suitable antioxidants are amino acids such as glycine, histidin, tyrosine, tryptophane, and derivatives thereof, imidazols such as urocaninic acid, and derivatives thereof, peptides, e.g. D,L-carnosin, D-carnosin, L-carnosin and their derivatives (e.g. anserin), carotinoids, carotins, (e.g. α-carotin, β-carotin, lycopin) and derivatives thereof, chlorogenic acid and

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their derivatives, liponic acids and their derivatives (e.g. dihydroliponic acid), aurothioglucose, propylthiouracil, and other thiols (e.g. thioredoxin, glutathion, cystein, cystin, cystamin, and glycosyl-, N-acetyl-, methyl-, ethyl-, propyl-, amyl-, butyl- and lauryl-, palmitoyl-, oleyl-, γ-linoleyl-, cholesteryl- and glyceryl-ester), as well as, salts thereof, dilaurylthiodipropionate distearylthiodipropionate thiodipropionic acid and derivatives thereof (esters, ether, peptide, lipide, nucleotide, nucleoside and salts), as well as, sulfoximins (e.g.buthioninisulfoximine, homocysteinsulfoximine, butioninsulfone-, penta-,hexa-, heptathioninsulfoximine) in very low tolerable concentrations (e.g. pmol to μmol/kg), further (metal)-chelators (e.g. α-hydroxy fatty acids, palmitine acids, phytine acids, lactoferrin), α-hydroxy acids, such as citric acid, lactic acid or malic acid, humic acid, bile acid, bile extracts, bilirubin, biliverdin, EDTA, EGTA and derivatives thereof, unsaturated fatty acids and derivatives thereof, such as y-linolenic acid, linoleic acid or oleic acid, foleic acid and derivatives thereof, ubiquinone, ubiquinol, and derivatives thereof, vitamin C and derivatives, such as ascorbylpalmitate, Mg-ascorbylphosphate or ascorbylacetate, tocopherols and derivatives, such as vitamin-E-acetate, vitamin A and derivatives, such as vitamin-A-palmitate, as well as $e^{-i\omega}$ coniferylbenzoate of benzoin resin, rutinic acid and derivatives thereof, α glycosylrutine, ferulic acid, furfurylidenglucitol, carnosin, butylhydroxytoluol, butvlhydroxyanisol, nordihydroguaiaretic acid resin, nordihydroguaiaretic acid, trihydroxybutyrophenon, uric acid and derivatives thereof, mannose and derivatives thereof, superoxide dismutase, zinc and derivatives thereof, such as ZnO and ZnSO₄, selenium and derivatives thereof, such as seleniummethionin, stilbenes and derivatives thereof, such as stilbeneoxide, trans stilbene oxide, and the suitable derivatives (salts, esters, ethers, sugars, nucleotides, nucleosides, peptides, lipids) of the mentioned agents.

In order to improve flowability it is possible to add hydrotropes, such as ethanol, isopropanol, or polyols. Preferred polyols have 2 to 15 carbon atoms and at least 2 hydroxyl groups. The polyols may have additional functional groups, in particular amino groups, respectively they may be modified with nitrogen. Typical examples are:

- glycerin;
- alkyleneglycols, such as ethyleneglycol, diethyleneglycol, propylene glycol, butylene glycol, hexylene glycol as well as

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- polyethyleneglycols with a average molecular weight of 100 to 1000 Dalton;
- technical oligoglycerin mixtures with a self condensation degree of 1.5 to 10 such as technical diglycerin mixtures with a diglycerin concentration of 40 to 50 % w/w;
- methyol compounds, in particular trimethylolethane, trimethylolpropane, trimethylolbutane, pentaerythrite and dipentaerythrite;
- lower alkylglucosides, in particular those having 1 to 8 carbon atoms in alkyl group, such as methyl- and butylglucoside
- sugar alcohols having 5 tot 12 carbon atoms, such as sorbitol or mannitol;
- sugars having 5 to 12 carbon atoms, such as glucose or saccharose
- amino sugars, such as glucamine;
- dialcoholamines such as diethanolamine or 2-amino-1,3-propanediol.

Examples of suitable preservatives are phenoxyethanol, formaldehyde, parabene, pentadiol or sorbic acid, as well as compounds mentioned in "Anlage 6, Teil A und B der Kosmetikverordnung". Examples of suitable insect-repellents are N,N-diethyl-m-toluamide, 1,2-pentadiol or ehtyl butylacetylaminoproprionate. Dihydroxy-aceton is a suitable tanning agent. Suitable tyrosine inhibitors - which prevent the formation of melanin and are employed in depigmentation agents - are for example arbutin, koji acid, coumarinic acid and ascorbic acid (vitamin C).

Suitable perfume oils are mixtures of natural and synthetic fragrant compounds. Natural fragrant compounds include extracts of flowers, such as lily, lavender, rose, jasmine, neroli and yiang-yiang, stems and leafs such as geranium, patchouli, petit-grain, fruits such as aniseed, coriander, cumin, juniper, fruit skins or peels, such as those of bergamot, lemons, oranges, roots such as macis, angelica, celery, cardamom, costus, iris, calmus, woods such as pine, sandal, guaja, cider and rose, herbs and grasses such as, estragon, lemon grass, sage, thyme, needles and branches, grove, spruce, pine, larch, resins and balms, such as galbanum, elimi, benzoe, myrrh, olibanum, opoponax.

Furthermore materials of animal origin can be used such as civet and

castoreum. Typical synthetic fragrant compounds are esters, ethers,

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aldehydes, ketones, alcohols and hydrocarbons. Examples of suitable esters are benzylacetate, phenoxyethylisobutyrate, p-tert.-butylcyclohexylacetate, linalylacetate, dimethylbenzylcarbinylacetate, phenylethylacetate, linalylbenzoate, benzylformiate, ethylmethylphenylglycinate, allylcyclohexylpropionate, styrallylproprionate and benzylsalicylate. Examples 5 of suitable ethers are benzylethyl sethers, examples of suitable aldehydes are linear alkanals having 8 to 18 carbon atoms, citral, citronellal, citronellyloxyacetaldehyde, cyclamealdehyde, hydroxycitronellal, lilial and bourgeonal, examples of suitable ketones are jojones, α-isomethylionon and methylcedrylketon, Examples of suitable alcohols are anethol, citronellol, 10 eugenol, iso-eugenol, geraniol, linalool, phenylethylalcohol and terpineol. Primary examples of hydrocarbons are terpenes and balms. Preferably however, mixtures of several fragrant compounds are employed, together resulting in a preferred aroma. In addition, etheric oils of low volatility, are suitable perfume oils. Examples of these are sage oil, camille oil, clove oil, 15 balm mint oil, mint oil, cinnamon oil, linden-blossom oil, juniper oil, vetiver oil, olibanum oil, galbanum oil, labolanum oil and lavender oil. Preferred are bergamot oil, dihydromyrcenol, lilial, lyral, citronellol, phenylethyl alcohol, αhexylcinnamon aldhyde, geraniol, benzylacetone, cyclamen aldehyde, linalool, boisambrene forte, ambroxan, indol, hedione, sandelice, citron-oil, mandarin-20 oil, orange-oil, allylamylglycolate, cyclovertal, lavender-oil, muscate sage-oil, B-damascone, geranium-oil bourbon, cyclohexylsalicylate, vertofix coeur, iso-Esuper, fixolide NP, evemyl, iraldein gamma, phenyl-acetic acid, geranyl acetate, benzyl acetate, rose-oxide, romilate, irotyl and floramate, either employed alone or in a mixture. 25

Suitable colorants are any colorants that are suitable for cosmetic purposes, as for example mentioned in the publication "Kosmetische Farbemittel" der Farbstoff-kommission der Deutschen Forschungsgemeinschaft, Verlag Chemie, Weinheim, 1984, S. 81-16". Such colorants are usually employed in concentrations varying from 0.001 to 0.1 % w/w based on the weight of the composition.

The total amount of additives can vary from 1 to 50 preferably from 5 to 40 % w/w based on the weight of the composition.

The balance of the composition will generally be made up by water. Optionally, a small amount of an alcohol, such as ethanol or isopropanol may be present, e.g. to achieve a disinfecting effect. Water will typically be present

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in an amount ranging from 50 to 95 wt.%, based on the weight of the composition.

The compositions can be prepared according to the usual cool or heated processes; a preferred method of preparation is a phase-inversion temperature method.

Dependent on the chosen ingredients of the composition as set forth above, a cosmetics composition according to the invention may find application as a sun cream or lotion, body milk, shampoo, bathing or shower gel, hair care product, deodorant or moisturizing cream. If desired, the present composition may also be employed in a pharmaceutical setting, for instance as an ointment. In such a case, the composition will further comprise a pharmaceutically active agent or a bioactive agent.

The invention will now be elucidated by the following, non-restrictive examples.

Example 1

A composition was prepared of the following ingredients in the following amounts (wt.%):

20	- Surfactants:	Magnesium Laureth Sulfate (1)	11.43
		Lauryl Glucoside (2)	5.19
	- pH regulator: Citric acid		0.11
	- Preservative: Kathon CG		0.06
	- Thickener: Xant	0.80	
25	- Moisturizing agent: Glycerin		5.00
	- Emollient: Cetiol HE		2.00
	- Conditioning agent: Merquat Plus 3331		1.00
	- Perfume		1.00
	- Coloring agent:	Patentblue V E 131	0.0015
30	- Water		Balance

The composition was prepared by first adding the water to a vessel. Next, in subsequent order, the preserative and the thickener were added. These components were mixed and homogenized until the thickener was swollen and fully dispersed. To the obtained dispersion, the surfactants were added separately with mixing to fully disperse the surfactant after each

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addition. The remaining ingredients, except the citric acid, were then added and mixing was continued until all were fully dispersed. Finally, the pH was adjusted by addition of the citric acid.

This composition was then cooled to below 10°C. The propellant to be added, isopentane, was also cooled to said temperature. The composition and the propellant were mixed with one another while taking care that no air was incorporated at constant temperature. The propellant was added in an amount to finally reach a concentration of 6 wt.%, with respect to the total weight of the final composition. After thorough mixing, the composition was allowed to warm up and brought into a suitable plastic container while still having a temperature below 20°C.

Example 2

Six compositions were prepared. Three were in accordance with the present invention (A, B, C), and three were not (CA, CB, CC). For the preparation of the compositions, a surfactant mixture comprising a thickener was mixed with pentane at a pH of 5.5-6.5. The viscosities were measured using the Brookfield method at 23°C ('Spindel TE', 5 rpm) in mPas. The results are shown in Table 1.

Table 1: Gels of surfactant mixtures (amounts in weight

percentages) CB Composition В \mathbf{C} CA \cdot CC Texapon® N70 8 8 4 8 8 4 Sodium Laureth Sulfate Plantacare® 818 1 1 1 1 Coco-Glucosid Dehyton® K 1 1 1 1 1 1 Cocoamidopropylbetain Glyceryl-PEG-140-4 4 4 4 4 4 tristearate Pentane 6 6 6 Water ad 100 Viscosity - 1 hour 30,500 25,000 34,000 3000 3200 2900 - 4 weeks-29,600 27,000 34,500 2700 2600 2800

Int. pat. appln. no. PCT/NL00/00599 Our letter of October 15, 2001M JC10 Rec'd PCT/PTO 1 2 FEB 2002

Amended Claims

- 1. Cosmetics composition comprising a thickener, isopentane as propellant, a surfactant and water, wherein the composition is contained in a container under a pressure of at least 0.1 bar in excess of atmospheric pressure and below 3 bar.
- 5 2. Composition according to claim 1, wherein the pressure is no more than 2 bar.
 - 3. Composition according to claim 1 or 2, wherein the container is a plastic container.
- 4. Composition according to any of the claims 1-3, wherein said composition comprises at least two surfactants and a hydrophobic compound having a HLB-value of less than 10 wherein said composition further has a viscosity according to Brookfield (23°C, Spindle TE, 5 RPM) of in the range of 5,000 to 50,000 mPas.
- 5. Composition according to any of the preceding claims, comprising •
 15 from 0.01 to 30 wt.% of thickener, from 1 to 15 wt.% of propellant, from 0.5 to
 50 wt.% of surfactant and the balance being water and other customary body
 care ingredients.
 - 6. Composition according to any of the preceding claims, wherein the thickener is chosen from the group of gums, poly(meth)acrylates, polymers based upon aerosil-types, polysaccharides, high molecular polyethyleneglycolmono- and diesters of fatty acids, polyacrylamides, polyvinylalcohols, polyvinylpyrrolidons, esters of fatty acids with polyols, fatty alcoholethoxylates, alkyloligoglucosides and sugar-esters.
- 7. Composition according to claim 6, wherein the thickener is chosen from xanthan gom, guar-guar, agar-agar alginates, tyloses, carboxymethylcellulose, hydroxyethylcellulose.

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- 8. Composition according to any of the preceding claims further comprising one or more ingredients chosen from the group of pH regulating agents, oil bodies, emulsifying agents, preservatives, perfumes, moisturizing agents, UV-filters, emollients, superfatting agents, brighteners, strength improving agents, silicon agents, fats, waxes, lecithins, phospholipids, stabilizing agents, anti-bacterial agents and other bioactive agents, odorabsorbing agents, antiperspirants, antidandruff agents, film-forming agents, ewelling agents, antioxidants, insect-repellents, hydrotropes, tanning agents, tyrosin inhibitors, solubilizers and colorants.
- 9. Composition according to any of the preceding claims, wherein said composition comprises a fatty alcohol preferably of the formula R₁OH, R₁ being a aliphatic hydrocarbon group containing 6 to 22 carbon atoms and 0,1,2, or 3 double bonds.
 - 10. Composition according to any of the preceding claims, wherein said composition comprises a fatty alcoholalkoxylate preferably of the formula R₂O(AlkO)_mH, R₂ being an aliphatic hydrocarbon group containing 6 to 22 carbon atoms, m being an integer from 1 to 30 and AlkO being an alkyleneoxide.
- 11. Composition according to any of the preceding claims, wherein said composition comprises a fatty alcoholalkoxylate of the formula R₂O(AlkO)_mH, R₂ being an aliphatic hydrocarbon group containing 8 to 22 carbon atoms, m being an integer from 5 to 20 and AlkO being chosen from ethyleneoxide and propylene oxide.
 - 12. Composition according to any of the claim 4 to 11, wherein the viscosity according to Brookfield (23°C, Spindle TE, 5 RPM) of said composition is in the range of 10,000 to 50,000, preferably in the range of 20,000 to 30,000 mPas.
 - 13. Composition according to any of the preceding claims wherein said composition is a gel.

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- 14. Composition according to any of the preceding claims wherein said composition comprises:
 - a) 0.01 to 30 % w/w of a thickener,
 - b) 0.1 to 20 % w/w of a hydrophobic compound having an HLB value of less than 10,
 - c) 0.5 to 40 % anionic surfactants,
 - d) 0.25 to 5 % amphotoric surfactants, and/or
 - e) 0.5 to 40 % nonionic surfactants,
- and is further characterized by the composition having a viscosity of 5,000 to 50,000 mPas and the weight ratio of components c:d or c:e being in the range of 2:1 to 8:1.
 - 15. Container comprising a cosmetics composition according to any of the preceding claims.
- 15 16. Use of a composition according to any of the claims claim 1-14 as a sun cream or lotion, body milk, shampoo, bathing or shower gel, ointment, deodorant, hair care product or moisturizing cream.
 - 17. Use of isopentane as a propellant incorporated into a cosmetics composition.

(19) World Intellectual Property Organization International Bureau



(43) International Publication Date 8 March 2001 (08.03.2001)

PCT

(10) International Publication Number WO 01/15659 A3

(51) International Patent Classification⁷: 7/16, 7/15, 7/11, 7/06, 7/00

A61K 7/32,

- (21) International Application Number: PCT/NL00/00599
- (22) International Filing Date: 29 August 2000 (29.08.2000)
- (25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

199 41 933.7 00200043.8 30 August 1999 (30.08.1999) DE 6 January 2000 (06.01.2000) EP

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- (84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

- With international search report.
- (88) Date of publication of the international search report: 25 May 2001

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

/15659 A

(54) Title: AQUEOUS LOW PRESSURE COSMETIC COMPOSITION COMPRISING A PROPELLANT, A THICKENER AND A SURFACTANT

Declaration and Power of Attorney Patent Application (Design or Utility)

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name,

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled: "Cosmetics composition".

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	is attached hereto		
図	is attached hereto was filed on February 12, 2002, as appl	cation serial no. 10/049,895 and or	PCT
	International Application number PCT/N	L00/00599.and was amended on	(ìf
	applicable).	' -	

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

Lacknowledge the duty to disclose to the U.S. Patent and Trademark Office all information know to me to be material to patentability as defined in 37 C.F.R.§1.56.

I hereby claim foreign priority benefits under 35 U.S.C.§119(a)-(d) or 35 U.S.C.§365(b) of any foreign application(s) for patent or inventor's certificate, or 35 U.S.C.§365(a) of any PCT International application which designated at least one country other than the United States, listed below and have also identified below any foreign application for patent or inventor's certificate of PCT International application having a filing date before that of the application on which priority is claimed.

	Prior Foreign Application(s)		
Country	Day/Month/Year Filed 30 August 1999		
Country	Day/Month/Year Filed		
	6 January 2000 Day/Month/Year Filed		
	DE		

I hereby claim the benefit under 35 U.S.C. §119(e) of any United States provisional application(s) listed below:

Prior Provisional Application(s)		
Serial Number	Day/Month/Year Filing Date	
Serial Number	Day/Month/Year Filing Date	
Serial Number	Day/Month/Year Filing Date	
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I hereby claim the benefit under 35 U.S.C. §120 of any United States application(s), or under 35 U.S.C. §365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of 35 U.S.C. §112, I acknowledge the duty to disclose to the U.S. Patent and Trademark Office all information known to me to be material to patentability as defined in 37 C.F.R.§1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application:

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Serial Number	Day/Month/Year Filed	Status (petented, pending, abandoned)	
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Serial Number	Day/Month/Year Filed	Status (patented, pending, abendoned)	

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C.§1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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Signature of Inventor		Date

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the specification of which

X .	is attached hereto was filed on February 12, 2002, as a International Application number PC	ipplication serial no. 10/049,895 and or T/NL00/00599 and was amended on	PCT (if
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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C.§1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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Jon R. Trembath		45,924
Ido Tuchman		42,189
Marcia A. Tunheim		27,403
Albert L. Underhill		32,179
J. Derek Vandenburgh		33,044
John R. Wahl		43,245
Karrie G. Weaver	•	20,890
Paul A. Welter		43,261
Brian Whipps John E. Whitaker		42,222
		41,376
J. Scott Wickhem		27,054
Douglas J. Williams James D. Withers		40,376
James D. Withers Jonelie Witt		41,980
Tong Wu		43,361
Min S. Xu	<i>)</i> .	39,536
Anthony R. Zeuli		45,255

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case to them and by whom/which I hereby declare that I have consented after full disclosure to be represented unless/until I instructed otherwise.

Please direct all correspondence in this case to at the address indicated below:

MERCHANT & GOULD P.C. P.O. Box 2903 Minneapolis, Minnesota 55402-0903 United States of America

Family Name	First Given Name	Second Given Name
Voss	Eckart	Karl Heinz
	Residence and Citizenship	
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Den Haag	the Netherlands	Germany
	Post Office Address	
Street Address	City	State & Zip Code or Country
Sportlaan 157	Den Haag	2566 KA
Signature of Inventor		Date

Family Name	First Given Name	Second Given Name
Knebel	Silke	Katharina
,	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
Düsseldorf	Germany	Germany
	Post Office Address	l de la companya de l
Street Address -	City	State & Zip Code or Country
Am Brückerstrasse 534	Düsseldorf	D-40591
Signature of Inventor		Date

	Full Name of Third Inventor, if a	
Family Name	First Given Name	Second Given Name
Monreal	Michele	·
	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
Kurten	Germany	Germany
	Post Office Address	
Street Address	City	State & Zip Code or Country
Kölnerstrasse	Kurten	D-51515
Signature of Inventor		Date

	Full Name of Fourth Inventor, if ar	ny
Family Name	First Given Name	Second Given Name
Withell	Trevor	Keith
	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
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	Post Office Address	
Street Address	City	State & Zip Code or Country
Drogetsberg 12 1144 Aurali	Düsseldorf Shin Cham	D-40489 RAZ+ SUP
Signature of Inventor	3600	Date 105 60 02

Family Name	Full Name of Fifth Inventor, if a	Second Given Name
***	Gerrit	
Nieman		
	Residence and Citizenship	
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Zoetermeer	the Netherlands	the Netherlands
	Post Office Address	
Street Address	City	State & Zip Code or Country
Lijnbaan 241	Zoetermeer	2728 AE
Signature of Inventor	,	Date
_		

Family Name	Full Name of Sixth Inventor, if a	Second Given Name
Hensen	Herman	
Name of the second seco	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
Haan	Germany	Germany
	Post Office Address	
Street Address	City	State & Zip Code or Country
Rathmacherweg 13	Haan	D-42781
Signature of Inventor		Date

Schmiedel Peter Residence and Citizenship City of Residence Düsseldorf State or Country of Residence Germany Germany Post Office Address Street Address City State & Zip Code or Country Hassiestrasse 62 Signature of Inventor Date	Residence and Citizenship City of Residence State or Country of Residence Country of Citize Düsseldorf Germany Germany	shin
Residence and Citizenship City of Residence Düsseldorf State or Country of Residence Germany Post Office Address Street Address City Hassiestrasse 62 City City City City City City City Cit	Residence and Citizenship City of Residence State or Country of Residence Country of Citize Düsseldorf Germany Germany	shin
Düsseldorf Germany Germany Post Office Address Street Address City State & Zip Code or Count Hassiestrasse 62 Düsseldorf D-40599	Düsseldorf Germany Germany	schin
Düsseldorf Germany Germany Post Office Address Street Address City State & Zip Code or Count Hassiestrasse 62 Düsseldorf D-40599	Düsseldorf Germany Germany	121116
Street Address Street Address City State & Zip Code or Count D-40599 D-40599		
Hassiestrasse 62 Düsseldorf D-40599	Fost Unice Address	
Hassiestrasse 62 Düsseldorf D-40599	Street Address City State & Zip Coo	e or Count

Ĭ,

Declaration and Power of Attorney Patent Application (Design or Utility)

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name,

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled: "Cosmetics composition".

the specification	on of which
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is attached hereto	₹	
was filed on February 12, 2002, as	application senal no. 10/049,895 and	or PCT
 International Application number PC	CT/NL00/00599 and was amended on	. (if
applicable).	1 .	

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose to the U.S. Patent and Trademark Office all information know to me to be material to patentability as defined in 37 C.F.R.§1.56.

I hereby claim foreign priority benefits under 35 U.S.C.§119(a)-(d) or 35 U.S.C.§365(b) of any foreign application(s) for patent or inventor's certificate, or 35 U.S.C.§365(a) of any PCT International application which designated at least one country other than the United States, listed below and have also identified below any foreign application for patent or inventor's certificate of PCT International application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application(s)		
Country	Day/Month/Year Filed 30 August 1999	
Country	Day/Month/Year Filed	
Country	6 January 2000 Day/Month/Year Filed	
	Country DE Country EP	

I hereby claim the benefit under 35 U.S.C. §119(e) of any United States provisional application(s) listed below:

Prior Provisional Application(s)		
Serial Number	Day/Month/Year Filing Date	
Serial Number	Day/Month/Year Filing Date	
Serial Number	Day/Month/Year Filing Date	
<u> </u>	· · · · · · · · · · · · · · · · · · ·	

I hereby claim the benefit under 35 U.S.C. §120 of any United States application(s), or under 35 U.S.C. §365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of 35 U.S.C. §112, I acknowledge the duty to disclose to the U.S. Patent and Trademark Office all information known to me to be material to patentability as defined in 37 C.F.R.§1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application:

Prior U.S. or International Application(s)		
Serial Number	Day/Month/Year Fifed	Status (patented, pending, abandoned)
Serial Number	Day/Month/Year Filed	Status (patented, pending, abandoned)
Serial Number	Day/Month/Year Filed	Status (patented, pending, abandoned)

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C.§1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Power of Attorney

As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith.

/	
Attorney	Registration Number
John W. Albrecht	40,481
M. Jeffer All	46,359
Gregg I. Anderson	28,828
Brian H. Batzli	32,960
John L. Beard	27,612
John M. Berns	43,496
Bruce E. Black	41,622
John W. Branch	41,633
Dennis C. Bremer	40,528
Steven C. Bruess	3 4,13 0
Linda M. Byrne	32,404
Keith Campbell	P-46,597
Alan G. Carlson	25,959
Philip P. Caspers	33,227
James R. Chiapetta	· 39,634
John A. Clifford	30,247
Richard J. Coldren	44,084
Ronald A. Daignault	25,968
Dennis R. Daley	34,994
Leslie E. Dalglish	40,579
Juie R. Daulton	36,414
Katherine M. DeVries Smith	42,157
Mark J. DiPietro	28,707
Robert T. Edell	^20,187
Sandra Epp Ryan	39,667
Robert J. Glance	40,620
Matthew J. Goggin	44,125
Charles E. Golla	26,896
Alan G. Gorman	38,472
John D. Gould	18,223
Richard Gregson	41,084
John J. Gresens	ິ33,112 46,754
Samuel A. Hamer	29,165
Curtis B. Hamre	P-46,759
Kevin C. Harrison	42,660
Brett A. Hertzberg	31,838
Randall A. Hillson	42,668
Richard J. Holzer, Jr. Scott W. Johnston	39,721
Natalie D. Kadievitch	34,196
Shaukat Karjeker	34,049
Joseph M. Kastelic	37,160
Denise Kettelberger	33,924
Jeramie J., Keys	42,724
Homer L. Knearl	21,197
Alan W. Kowalchyk	31,535
A Marie Val a Se statistical	•

Katherine M. Kowalchyk		36,848
Paul E. Lacy		38,946
James A. Larson		40,443
Andrew J. Leon		46,869
Christopher J. Leonard		41,940
Mara E. Liepa		40,066
Timothy A. Lindquist		40,701
Lawrence E. Lycke		38,540
Denise L. Mayfield		33,732
Daniel W. McDonald		32,044
William F. McIntyre, Jr.		44,921
M. Todd Mitchem		40,731
Douglas P. Mueller	· ·	30,300
A. Shane Nichols		43,836
Daniel M. Pauly		40,123
Bryan K. Phillips		P-46,990
John B. Phillips	<u> </u>	37,206
Paul Prendergast	•	46,068
Melissa J. Pytel		41,512
Terry Qualey		25,148
John C. Reich	1 7	37,703
Earl D. Reiland		25,767
Lisa A. Samuels		43,080
David G. Schmaltz		39,828
Mark D. Schuman		31,197
Micheal D. Schumann		30,422
Timothy B. Scull		42,137
Gregory A. Sebald		33,280
Mark T. Skoog		40,178
Steven J. Spellman		45,124
Kirstin L. Stoll-DeBell		43,164
John P. Sumner		29,114
Erik G. Swenson		45,147
David K. Tellekson		32,314
Jon R. Trembath		38,344
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Paul A. Welter		20,890
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John E. Whitaker		42,222
J. Scott Wickhem	-	41,376
Douglas J. Williams		27,054
James D. Withers		40,376
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Tong Wu		43,361
Min S. Xu		39,536
Anthony R. Zeuli		45,255

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MERCHANT & GOULD P.C. P.O. Box 2903 Minneapolls, Minnesota 55402-0903 United States of America

Family Name	First Given Name	Second Given Name
Voss	Eckart	Karl Heinz
	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
Den Haag	the Netherlands	Germany
	Post Office Address	
Street Address	City	State & Zip Code or Country
Sportlaan 157	Den Haag	2566 KA
Signature of Inventor		Date

	Full Name of Second Inventor, if	any
Family Name	First Given Name	Second Given Name
Knebel	Silke	Katharina
	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
Düsseldorf	Germany	Germany
	Post Office Address	
Street Address	City	State & Zip Code or Country
Am Brückerstrasse 534	Düsseldorf	D-40591
Signature of Inventor	1.	Date

Family Name	First Given Name	Second Given Name
Monreal	Michele	
	Residence and Citizenship	
City of Residence	State of Country of Residence	Country of Citizenship
Kurten	Germany	Germany
	Post Office Address	
Street Address	City	State & Zip Code or Country
Kölnerstrasse	Kurten	D-51515
Signature of Inventor		Date

Full Name of Fourth Inventor, If any			
Family Name	First Given Name	Second Given Name	
Withell	Trevor	Keith	
	Residence and Citizenship		
City of Residence	State or Country of Residence	Country of Citizenship	
Düsseldorf	Germany	Great Brittain	
	Post Office Address		
Street Address	City	State & Zip Code or Country	
Drögelsberg	Düsseldorf	D-40489	
Signature of Inventor		Date	

Will the second second

Family Name	First Given Name	Second Given Name
Nieman	Gerrit	Good at Given Name
	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
Zoetermeer	the Netherlands	the Netherlands
	Post Office Address	
Street Address	City	State & Zip Code or Country
Lijnbaan 241	Zoetermeer	2728 AE
Signature of Inventor		Date

	Full Name of Sixth Inventor, if a	
Family Name	First Given Name	Second Given Name
Hensen	Herman	
	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
Haan	Germany	Germany
	Post Office Address	
Street Address	City	State & Zip Code or Country
Rathmacherweg 13	Haan	D-42781
Signature of Inventor //	77	Date
1/2	ricum Hirsen	13 9.02

Family Name	Full Name of Seventh Inventor, if First Given Name	Second Given Name
Schmiedel	Peter	
	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
Düsseldorf	Germany	Germany
	Post Office Address	
Street Address	City	State & Zip Code or Country
Hasslestrasse 62	Düsseldorf	D-40599
Signature of Inventor	Time and the second	Date 77 Cut OP

Declaration and Power of Attorney Patent Application (Design or Utility)

As a below named inventor, I hereby declare that:

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the	specification of which		
	is attached hereto was filed on February 12, 2002, as a International Application number PC		
	applicable)."	•	

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Prior Foreign Application(s)		
Number 199 41 933.7	Country	Day/Month/Year Filed 30 August 1999
Number 00200043.8	Country	Day/Month/Year Filed 6 January 2000
Number	Country	Day/Month/Year Filed
Number	Country	pay/Monuti Year Filed

I hereby claim the benefit under 35 U.S.C. §119(e) of any United States provisional application(s) listed below:

Prior Provisional Application(s)		
Serial Number Day/Month/Year Filing Date		
Serial Number	Day/Month/Year Filing Date	
Serial Number	Day/Month/Year Filing Date	
<u> </u>		

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Serial Number	Day/Month/Year Filed	Status (patented, pending, abandoned)
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M. Jeffer Ali	46,359
Gregg I. Anderson	28,828
Brian H. Batzli	32,960
John L. Beard	27,612
John M. Berns	43,496
Bruce E. Black	41,622
John W. Branch	41,633
Dennis C. Bremer	40,528
Steven C. Bruess	3 4 ,130
Linda M. Byrne	32,404
Keith Campbell	P-46,597 25,959
Alan G. Carlson	25,959
Philip P. Caspers	33 227
James R. Chiapetta	39,634
John A. Clifford	30,247
Richard J. Coldren	44,084
Ronald A. Daignault	25,968
Dennis R. Daley	34,994
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Juie R. Daulton	36,414
Katherine M. DeVries Smith	42,157
Mark J. DiPietro	28,707
Robert T. Edell	20,187
Sandra Epp Ryan	39,667
Robert J. Glance	40,620
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Alan G. Gorman	38,472
John D. Gould	18,223
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Curtis B. Hamre	29,165
Kevin C. Harrison	P-46,759
Brett A. Hertzberg	42,660
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Joseph M. Kastelic	37,160
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	•

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Christopher J. Leonard		41,940
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Timothy A. Lindquist		40,701
Lawrence E. Lycke		38,540
Denise L. Mayfield	· .	33,732
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M. Todd Mitchem		40,731
Douglas P. Mueller		30,300
A. Shane Nichols		43,836
Daniel M. Pauly		40,123
Bryan K. Phillips		P-46,990
John B. Phillips		37,206
Paul Prendergast	•	46,068
Melissa J. Pytel		41,512
Terry Qualey		25,148
John C. Reich		37,703
Earl D. Reiland		25,767
Lisa A. Samuels		43,080
David G. Schmaltz		39,828
Mark D. Schuman		31,197
Micheal D. Schumann		30,422
Timothy B. Scull		42,137
Gregory A. Sebald		33,280
Mark T. Skoog		40,178
Steven J. Spellman		45,124
Kirstin L. Stoll-DeBell		43,164
John P. Sumner		29,114
Erik G. Swenson		45,147
David K. Tellekson	·	32,314
Jon R. Trembath	•	38,344
ldo Tuchman		45,924
Marcia A. Tunheim		42,189
Albert L. Underhill		27,403
J. Derek Vandenburgh		32,179
John R. Wahl		33,044
Karrie G. Weaver		43,245
Paul A. Welter		20,890
Brian Whipps	• •	43,261
John E. Whitaker	•	42,222
 J. Scott Wickhem 		41,376
Douglas J. Williams		27,054
James D. Withers		40,376
Jonelle Witt		41,980
Tong Wu		43,361
Min S. Xu		39,536
Anthony R. Zeuli	•	45,255

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MERCHANT & GOULD P.C. P.O. Box 2903 Minneapolis, Minnesota 55402-0903 United States of America

Family Name	First Given Name	Second Given Name
Voss	Eckart	Karl Heinz
	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
Den Haag	the Netherlands ///	Germany
	Post Office Address	
Street Address	City	State & Zip Code or Country
Sportlaan 157	Den Haag	2566 KA
Signature of Inventor	70 1 1/11/	Date 21. 09. 02

20

	Full Name of Second Inventor, if	any
Family Name	First Given Name	Second Given Name
Knebel	Silke	Katharina
	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
Düsseldorf	Germany /	Germany
	Post Office Address	
Street Address -	City	State & Zip Code or Country
Am Brückerstrasse 534	Düsseldorf	D-40591
Signature of Inventor		Date

Family Name	First Given Name	Second Given Name
Monreal	Michele	,
	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
Kurten	Germany (/	Germany
	Post Office Address	
Street Address	City	State & ZIp Code or Countr
Kölnerstrasse	Kurten	D-51515
Signature of Inventor		Date

Full Name of Fourth Inventor, If any Second Given Name Family Name First Given Name Withell Keith Trevor Residence and Citizenship
State or Country of Besidence
Germany
Post Office Address City of Residence Country of Citizenship Düsseldorf Great Brittain Street Address City State & Zip Code or Country Drögelsberg Signature of Inventor Düsseldorf D-40489 Date

5d

Family Name	First Given Name	Second Given Name
Nieman	Gerrit	<u> </u>
	Residence and Citizenship	
City of Résidence	State or Country of Residence)	Country of Citizenship
Zoetermeer	the Netherlands ////	the Netherlands
	Post Office Address	
Street Address	City	State & Zip Code or Country
Lijnbaan 241	Zoetermeer	2728 AE
Signature of Inventor		Date

6-D-

	Full Name of Sixth Inventor, if a	iny
Family Name	First Given Name	Second Given Name
Hensen	Herman	<u> </u>
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-	Post Office Address	
Street Address	City	State & Zip Code or Country
Rathmacherweg 13	Haan	D-42781
Signature of Inventor		Date
,		1

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	Full Name of Seventh Inventor, if	any
Family Name	First Given Name	Second Given Name
Schmiedel	Peter	<u> </u>
	Residence and Citizenship	
City of Residence	State or Country of Residence	Country of Citizenship
Düsseldorf	Germany (/ 127)	Germany
	Post Office Address	
Street Address	City	State & Zip Code or Country
Hasslestrasse 62	Düsseldorf	D-40599
Signature of Inventor		Date